

Psychometric properties and feasibility of use of dementia specific quality of life instruments for use in care settings: a systematic review

Article (Published Version)

Hughes, Laura J, Farina, Nicolas, Page, Thomas E, Tabet, Naji and Banerjee, Sube (2019) Psychometric properties and feasibility of use of dementia specific quality of life instruments for use in care settings: a systematic review. *International Psychogeriatrics*. pp. 1-15. ISSN 1041-6102, 1741-203X

This version is available from Sussex Research Online: <http://sro.sussex.ac.uk/id/eprint/81118/>

This document is made available in accordance with publisher policies and may differ from the published version or from the version of record. If you wish to cite this item you are advised to consult the publisher's version. Please see the URL above for details on accessing the published version.

Copyright and reuse:

Sussex Research Online is a digital repository of the research output of the University.

Copyright and all moral rights to the version of the paper presented here belong to the individual author(s) and/or other copyright owners. To the extent reasonable and practicable, the material made available in SRO has been checked for eligibility before being made available.

Copies of full text items generally can be reproduced, displayed or performed and given to third parties in any format or medium for personal research or study, educational, or not-for-profit purposes without prior permission or charge, provided that the authors, title and full bibliographic details are credited, a hyperlink and/or URL is given for the original metadata page and the content is not changed in any way.

REVIEW

Psychometric properties and feasibility of use of dementia specific quality of life instruments for use in care settings: a systematic review

Laura J. Hughes,¹ Nicolas Farina,¹  Thomas E. Page,² Naji Tabet,¹ and Sube Banerjee¹

¹Centre for Dementia Studies, Brighton and Sussex Medical School, Brighton, UK

²School of Psychology, University of Kent, Canterbury, UK

ABSTRACT

Background: Over 400,000 people live in care home settings in the UK. One way of understanding and improving the quality of care provided is by measuring and understanding the quality of life (QoL) of those living in care homes. This review aimed to identify and examine the psychometric properties including feasibility of use of dementia-specific QoL measures developed or validated for use in care settings.

Design: Systematic review.

Methods: Instruments were identified using four electronic databases (PubMed, PsycINFO, Web of Science, and CINAHL) and lateral search techniques. Searches were conducted in January 2017. Studies which reported on the development and/or validation of dementia specific QoL instruments for use in care settings written in English were eligible for inclusion. The methodological quality of the studies was assessed using the COSMIN checklist. Feasibility was assessed using a checklist developed specifically for the review.

Results: Six hundred and sixteen articles were identified in the initial search. After de-duplication, screening and further lateral searches were performed, 25 studies reporting on 9 dementia-specific QoL instruments for use in care home settings were included in the review. Limited evidence was available on the psychometric properties of many instruments identified. Higher-quality instruments were not easily accessible or had low feasibility of use.

Conclusions: Few high-quality instruments of QoL validated for use in care home settings are readily or freely available. This review highlights the need to develop a well-validated measure of QoL for use within care homes that is also feasible and accessible.

Key words: dementia, quality of life, care settings, feasibility, psychometrics

Introduction

There are approximately 16,000 care homes in the UK providing care for an estimated 416,000 people (Care Quality Commission, 2017a; NIHR, 2016); in the US over 2 million individuals are cared for in approximately 45,000 facilities (Centre for Disease Control and Prevention, 2017). These numbers are expected to increase with an ageing population (Prince *et al.*, 2014). It is estimated that anywhere

between 40% and 86% of people living in care homes have a dementia, either diagnosed or undiagnosed, or some form of memory impairment (Jagger and Lindesay, 1997; Livingston *et al.*, 2017). The large numbers of people living in care homes has led to a need to understand the outcomes and experiences of care by residents in order to make improvements to care. The quality of care provided in care homes and its consistency has been questioned, with calls to increase the services provided to vulnerable adults (Care Quality Commission, 2017b; Department of Health, Prime Minister's Office, 2015). It has been argued that one way we may be able to better understand the outcomes of care in order to improve it is through the measurement and understanding of quality of life (QoL) (Black, 2013; Edelman *et al.*, 2005).

Correspondence should be addressed to: Laura Hughes, Centre for Dementia Studies, Trafford Centre, Brighton & Sussex Medical School, University of Sussex, Falmer, Brighton BN1 9RY, UK. Phone: 01273 877896. Email: L.J.Hughes@bsms.ac.uk. Received 15 Jun 2018; revision requested 06 Sep 2018; revised version received 10 Oct 2018; accepted 26 Nov 2018.

Due to a growing recognition of the importance of the outcomes of care as well as the process of care (Sloane *et al.*, 2005), there has been a growing emphasis on measuring QoL as a means of understanding and improving care in care settings. An increasing amount of research in care homes is being carried out with QoL as an important outcome for evaluating interventions (Aspden *et al.*, 2014). A number of dementia-specific QoL measures have been developed; however, most of these were developed and evaluated in community-dwelling populations, rather than in care homes (Bowling *et al.*, 2015).

Previous reviews have examined dementia-specific QoL instruments in general (Bowling *et al.*, 2015; Ready and Ott, 2003), and generic QoL measures available for use in care homes (Aspden *et al.*, 2014). However, generic QoL measures have limitations in capturing the experiences of people with dementia (Smith *et al.*, 2005). No previous review has focused on the usability of the instruments (e.g. availability, cost, training, and how easy is it to score and interpret) in care homes settings. Ultimately, the usability of the instrument is likely to dictate whether an instrument is used or not. We therefore aimed to add to previous reviews by carrying out a systematic review to identify and examine not only the psychometric properties but also the usability of disease-specific instruments that measure the QoL of people living in care homes.

Methods

The protocol for this review is registered in the International Prospective Register of Systematic Reviews (PROSPERO) – CRD42017046272.

Inclusion criteria

Studies were included in the review if they met the following criteria: 1) They described the development and/or evaluation of an instrument, or described the adaptation and evaluation of an existing instrument for a care home population; 2) the instrument was a dementia-specific QoL instrument, and studies evaluating only generic health-related QoL instruments in a care home population were excluded; 3) the study population included residents living in a care home (including nursing homes), and data on this group were presented separately from any others studied; and 4) studies were published in English. There were no exclusion criteria based on diagnosis, or non-diagnosis, of dementia.

Search strategy

Articles were identified from initial searches in four electronic databases: PubMed, PsycINFO, Web of

Science, and the Cumulative Index to Nursing and Allied Health Literature (CINAHL). All searches were conducted in January 2017. There were no restrictions on date of publication. The following four combined search terms were used: 1) quality of life OR QOL OR health related quality of life OR HRQOL OR HRQL, AND 2) dementia OR Alzheimer's, AND 3) residential facilities OR residential OR care institutions OR long-term care OR nursing homes OR care homes OR residential care homes, AND 4) measure development OR valid* OR reliab* OR accuracy OR feasibility OR scale. Lateral searches involved checking the references of included studies (snowballing), and further searching for identified measures on PubMed and Google search engine. Two independent reviewers (LH & NF) screened article titles and abstracts against the predefined inclusion criteria. Full text articles were sought for all relevant studies. Any disagreements regarding inclusion were resolved through discussion by the two reviewers.

Data extraction

Two reviewers (LH & TEP) independently extracted the following data from included full-texts: name of instrument, country, language of instrument, sample characteristics (i.e., age, gender, and dementia severity), study design, measurement domains, individual items, number of items, response format, and evidence of reliability and validity.

The reviewers also extracted the following data about the usability of each instrument through reading the identified articles, internet searches (Google.com, first 30 hits) and a measure specific database (Mapi; <https://mapi-trust.org/>). Data included:

- Existence of a dedicated website (Yes/No) – The accessibility of an online website or repository, which contains dedicated information about the instrument
- User guide available (Yes/No) – The availability of a user guide or instructions about how to use the instrument
- Cost (paid, free, or other) – The cost of using the questionnaire, broken down by user type when provided
- Training – Details about what training is available for the instrument, whether it is a prerequisite, and the costs involved
- Time to complete measure (minutes)
- Time period the instrument captures
- Need for specialist software for scoring and interpretation (Yes/No)
- Guidance available for scoring and interpretation (Yes/No)

“Not known” was used in instances where we were unable to identify measure specific information.

Quality assessment

The methodological quality of the studies was assessed using the Consensus-based Standards for the selection of health Measurement Instruments (COSMIN) checklist (Mokkink *et al.*, 2010). This is a standardized tool which assesses the measurement properties of health-related instruments across nine domains (internal consistency, reliability, measurement error, content validity [including face validity], construct validity [subdivided into structural validity, hypotheses testing, and cross-cultural validity], criterion validity and responsiveness) with each domain rated using 5–18 items. Each item is rated as “excellent”, “good”, “fair”, or “poor” quality. A methodological quality score for each measurement property is obtained by taking the lowest rating of any item in that box (“worst score counts”). Two independent reviewers (LH & TEP) assessed the methodological quality of the included studies using the checklist. Studies will not be excluded based on COSMIN scores. Any disagreements in scoring were resolved through discussion and advice from a third reviewer (NF).

Data synthesis

A narrative synthesis was adopted to assess the feasibility of the QoL measures for use in care homes. No formal frameworks or criteria exist for the assessment of the feasibility of using QoL measures in care homes. Therefore, a set of criteria identified by the researchers in consultation with care home staff as important for using QoL measures in this setting was created *a priori*, based upon the working structure and practices of care and nursing homes in the UK.

Results

Search results

Initial database searches identified 616 articles, of which 269 were removed after duplicate deletion. The titles and abstracts of 347 articles were screened resulting in the exclusion of 308 articles. Full text papers of the remaining 38 articles were sought. After reviewing full texts, 19 articles met the inclusion criteria. Figure 1 illustrates the process. Six additional articles were identified from lateral searches.

In total we included 25 studies in the systematic review; these reported on nine different QoL instruments for use in care settings. The measures were: the Dementia Quality of Life (DQoL) (Brod *et al.*, 1999) ($n = 4$), Quality of Life in Alzheimer's Disease (QOL-AD) (Logsdon *et al.*, 1999) ($n = 1$), Quality of Life in Alzheimer's Disease nursing home version (QOL-AD NH) (adapted by Edelman and Fulton,

unpublished work) ($n = 6$), QUALIDEM (Ettema *et al.*, 2007b) ($n = 7$), Quality of Life in Late-Stage Dementia (QUALID) ($n = 6$) (Weiner *et al.*, 1999), Dementia Care Mapping (DCM) (Bradford Dementia Group, 2005) ($n = 5$), Alzheimer Disease Related Quality of Life (ADRQL) (Rabins *et al.*, 1999) ($n = 3$), ADRQL revised (Kasper *et al.*, 2009) ($n = 1$), and the Quality of Life in Dementia (QOL-D) (Albert *et al.*, 1996) ($n = 1$).

Study and measure characteristics

The sample size of the studies ranged from 20 (Beer *et al.*, 2009) to 973 (Bouman *et al.*, 2011). Table 1 outlines the characteristics of included studies. The studies recruited participants from the United Kingdom ($n = 8$), the United States ($n = 2$) Netherlands ($n = 3$), Spain ($n = 4$), Germany ($n = 5$), Sweden ($n = 1$), and Norway ($n = 2$).

Of the nine instruments identified, three instruments contained self-report and a proxy report questions (QOL-AD, QOL-AD NH, and QOL-D), one instrument contained a self-report questions only (DQoL), four instruments contained a proxy report questions only (QUALIDEM, QUALID, ADRQL, and ADRQL revised), and one instrument consists of a proxy observation tool (DCM). The number of measurement domains ranged from 1 (QOL-AD, QOL-AD NH, and QUALID) to 9 (QUALIDEM 37 item version). Five instruments have a Likert scale response format (DQoL, QOL-AD, QOL-AD NH, QUALID, and DCM), two have a frequency scale format (e.g., never, sometimes, often) (QUALIDEM and QOL-D), and two instruments have a dichotomous response format (e.g., agree/disagree) (ADRQL and ADRQL revised). The number of items in each instrument ranged from 6 (affect subscale of QOL-D) to 40 (ADRQL revised). See Supplementary Table 1 for full details.

Six instruments were developed specifically for use in care settings (QUALID, QUALIDEM, DCM, ADRQL, ADRQL revised, and QoL-D), and one was adapted for use in care settings and evaluated (QOL-AD NH). The QOL-AD and DQoL were evaluated for use in care settings but were not developed or adapted for this purpose.

Psychometric properties

The instrument with the most comprehensive evaluation of psychometric properties across the nine domains of the COSMIN checklist was the QUALID with seven domains assessed in two separate studies. The DQoL, QOL-AD, QUALIDEM, and DCM had four domains assessed in at least one study.

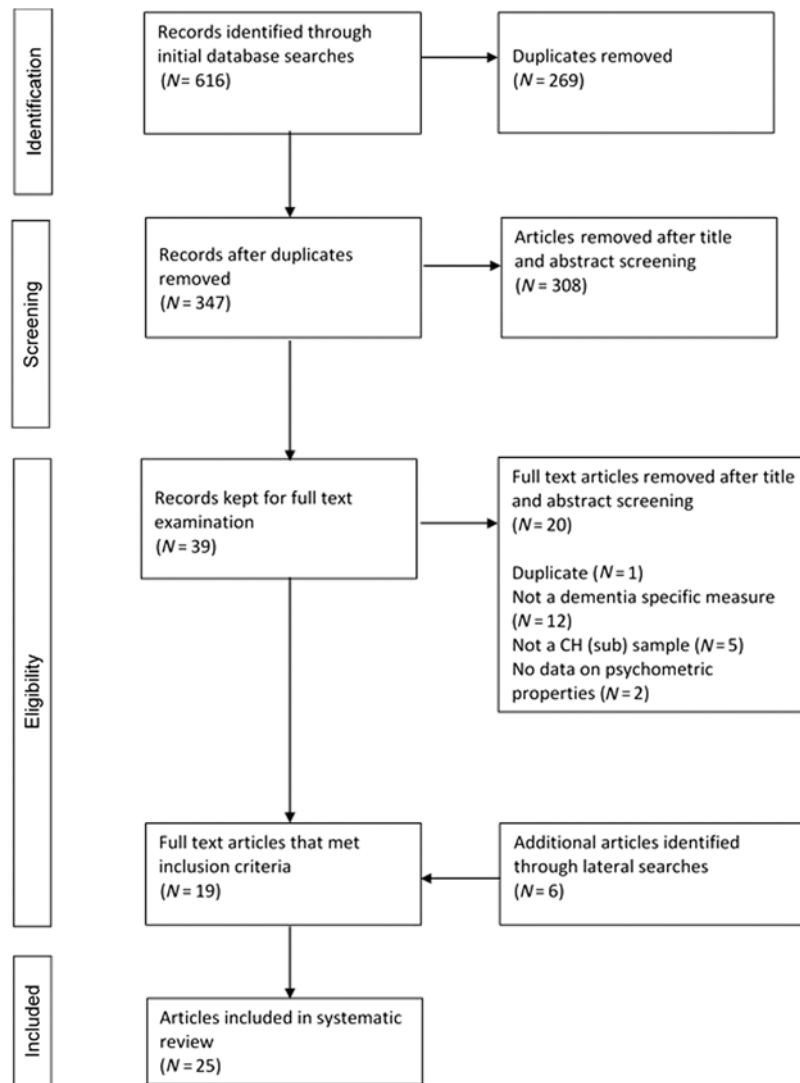


Figure 1. Flow diagram of study selection.

The instruments with the least domains assessed were QOL-AD NH ($n = 3$), QOL-D ($n = 3$), ADRQL ($n = 3$), and ADRQL revised ($n = 1$). See Table 2 for results of the COSMIN checklist.

Internal consistency

All nine instruments had internal consistency reported in at least one study. Instruments that were reported on the most were QUALID (six studies with poor to good ratings), and QUALIDEM (five studies, poor to excellent quality ratings). Most of the instruments had acceptable to good internal consistency across some studies. QOL-AD, ADRQL Revised, and QOL-AD NH all had the highest internal consistency scores in a single or small number of studies (all >0.80). QUALID had the most consistent high internal consistency scores for multiple studies (all >0.70). All other instruments had

variability in scores from poor to excellent internal consistency.

Reliability

Test retest and inter-rater reliability were rated in eight instruments ($n = 15$). The most assessed instruments were QUALIDEM, QUALID, and DCM ($n = 4$). QUALIDEM was rated fair to good, QUALID and DCM were both rated poor to fair. QOL-AD NH was assessed in three studies (poor to fair), DQoL was assessed in two (fair), and the QOL-AD (poor) ADRQL (fair), and QOL-D (fair) were assessed in one study. Time between tests for test-retest reliability ranged from 2–3 days (QUALID) to 12 months (DQoL). There was large variability in reliability scores for instruments. QOL-D and ADRQL both had the highest reliability, from a single study each (>0.95). QUALID had the highest

Table 1. Characteristics of included studies

	INSTRUMENT USED	COUNTRY	POPULATION CHARACTERISTICS (RESIDENTS)			
			N	GENDER (% F)	MEAN AGE (SD)	COGNITIVE IMPAIRMENT MEAN (SD)/N (%)
Adler and Resnick (2010)	DQOL	United Kingdom	486	80	83.8 (8.2)	MMSE 20.4 (5.3)
Beer <i>et al.</i> (2009)	QOL-AD NH	United Kingdom	20	70	87.5 (8.1)	MMSE 15.4 (5.2)
Bouman <i>et al.</i> (2011)	QUALIDEM	Netherlands	973	71–76 across four samples	83.9 (5.8)	GDS <7, N = 759 GDS 7, N = 214
Clare <i>et al.</i> (2014a)	QUALID	United Kingdom	105	69	81.47 (8.63)	FAST 6, N = 25 (24%) FAST 7, N = 80 (76%)
Crespo <i>et al.</i> (2011)	QOL-AD NH	Spain	209	79.4	86.07 (6.46)	MMSE 13.3 (5.88)
Dichter <i>et al.</i> (2011)	QUALIDEM	Germany	486	74 mild-moderate 79 severe-very severe	–	MMSE <24 = 15.4 (4) MMSE <10 = 3 (3)
Dichter <i>et al.</i> (2013)	QUALIDEM	Germany	634	78.6 mild-severe 77.7 very severe	85.8 (8.0) mild-severe 84.5 (8.2) very severe	FAST 2–6, N = 378 (100%) FAST 7, N = 256 (100%)
Dichter <i>et al.</i> (2014)	QUALIDEM	Germany	161	91 mild-severe 75 very severe	85.2 (7.0) mild-severe 84.3 (9.0) very severe	FAST 2–6, N = 108 (100%) FAST 7, N = 53 (100%)
Dichter <i>et al.</i> (2016a)	QUALIDEM	Germany	91	78 mild-severe 92 very severe	84.7 (8.3) mild-severe 84.1 (9.4) very severe	FAST 2–6, N = 55 (100%)
Dichter <i>et al.</i> (2016b)	QOL-AD NH	Germany	29	66	88.2 (7.1)	Not reported
Edelman <i>et al.</i> (2005)	QOL-AD NH DQoL ADRQL DCM	United Kingdom	172	83.7	85.8 (6.4)	MMSE 9.0 (6.9)
Ettema <i>et al.</i> (2007a)	QUALIDEM	Netherlands	202	–	–	GDS <7, N = 202 (100%)
Ettema <i>et al.</i> (2007b)	QUALIDEM	Netherlands*	238	76.5	84.69 (6.85)	GDS 2–6, N = 202 (100%) GDS 7, N = 36 (100%)
Falk <i>et al.</i> (2007)	QUALID	Sweden	169	88	86 (6.0)	MMSE 10 (7)
Fossey <i>et al.</i> (2002)	DCM	United Kingdom	123 (A) 54 (B)	76 cohort A 35 cohort B	82.3 (7.4) cohort A 78.2 (9.1) cohort B	Cohort A, MMSE 9.5 (7.7). CDR stages 1 = 19%, 2 = 38%, 3 = 43% Cohort B, 100% CDR stage 3
Garre-Olmo <i>et al.</i> (2010)	QUALID	Spain	160	76.9	82.6 (11.6)	MMSE = 4.1 (6.3)
Kasper <i>et al.</i> (2009)	ADRQL ADRQL revised	United States	89	–	– (–)	–
León-Salas <i>et al.</i> (2011)	QOL-AD	Spain	101	88.1	83.2 (6.3)	MMSE = 7.2 (6.1)

Table 1. Continued

	INSTRUMENT USED	COUNTRY	POPULATION CHARACTERISTICS (RESIDENTS)			
			N	GENDER (% F)	MEAN AGE (SD)	COGNITIVE IMPAIRMENT MEAN (SD)/N (%)
Mjorud <i>et al.</i> (2014)	QUALID	Norway	661	71.4	85.3 (8.6)	CDR stages 1: N = 147, 2: N = 222, 3: N = 290
Moyle <i>et al.</i> (2012)	DQoL QOL-AD NH	United Kingdom	61	70.5		MMSE 20–30 = 38 (62.3%) 10–19 = 22 (36.1%) 0–9 = 1 (1.6%)
Røen <i>et al.</i> (2015)	QUALID	Norway	169	56.6 reliability study 69.2 validity study	83.2 (7.5) reliability study 84.9 (6.7) validity study	MMSE 13.6 (6.1) reliability study MMSE 14.0 (6.0) validity study
Sloane <i>et al.</i> (2005)	QOL-D QOL-AD NH ADRQL DQoL DCM	United Kingdom	421	79	85 (–)	MMSE Mild <17 = 12% Moderate 10–16 = 25% Severe 3–9 = 27% Very severe <2 = 37%
Sloane <i>et al.</i> (2007)	DCM	United Kingdom	–	–	–	–
Villar <i>et al.</i> (2015)	DCM	Spain	68	66.7	86.08 (7.04)	MMSE = 18.09 (3.75)
Weiner <i>et al.</i> (1999)	QUALID	United States	42	67	80.6 (–)	MMSE = 11.53 (6.23)

*English language instrument used.

ADRQL, Alzheimer Disease Related Quality of Life; ADRQL revised, Alzheimer Disease Related Quality of Life revised version; DCM, Dementia Care Mapping; DQoL, Dementia quality of life; QOL-AD, Quality of Life in Alzheimer's Disease; QOL-AD NH, Quality of Life in Alzheimer's Disease nursing home version; QOL-D, Quality of Life in Dementia; QUALID, The quality of life in late-stage dementia; QUALIDEM, Quality of life instrument for proxy completion; MMSE, Mini Mental State Examination (Folstein *et al.*, 1975); FAST, Functional Assessment Staging (Reisberg, 1988); GDS, Global Deterioration Scale (Reisberg *et al.*, 1982); CDR, Clinical Dementia Rating Scale (Morris, 1997); – = Not reported.

Table 2. Results of consensus-based standards for the selection of health measurement instruments (COSMIN) checklist

QOL INSTRUMENT	AUTHOR(S)	INTERNAL CONSISTENCY	RELIABILITY	MEASUREMENT ERROR	CONTENT VALIDITY	STRUCTURAL VALIDITY	HYPOTHESIS TESTING	CROSS-CULTURAL VALIDITY	CRITERION VALIDITY	RESPONSIVENESS
DQoL	Edelman <i>et al.</i> (2005)	Good	•	•	•	Poor	Poor	•	•	•
	Adler and Resnick (2010)	Poor	Fair	•	•	Fair	•	•	Poor	•
	Moyle <i>et al.</i> (2012)	Poor	•	•	•	•	Fair	•	•	•
	Sloane <i>et al.</i> (2005)	Poor	Fair	•	•	•	Poor	•	•	•
QOL-AD NH	Edelman <i>et al.</i> (2005)	Good	•	•	•	Poor	Poor	•	•	•
	Moyle <i>et al.</i> (2012)	Poor	•	•	•	•	Fair	•	•	•
	Sloane <i>et al.</i> (2005)	Poor	Fair	•	•	•	Poor	•	•	•
	Crespo <i>et al.</i> (2011)	Poor	Fair	•	•	•	•	Poor	•	•
	Dichter <i>et al.</i> (2016b)	•	•	•	•	•	•	Fair	•	•
	Beer <i>et al.</i> (2009)	•	Poor	•	•	•	•	•	•	•
	Dichter <i>et al.</i> (2013)	Excellent	•	•	•	Excellent	•	•	•	•
QUALIDEM	Bouman <i>et al.</i> (2011)	Excellent	Fair	•	•	Excellent	•	•	•	•
	Dichter <i>et al.</i> (2011)	Good	•	•	•	Fair	•	Poor	•	•
	Ettema <i>et al.</i> (2007b)	Poor	Good	•	Fair	Good	•	•	•	•
	Dichter <i>et al.</i> (2016a)	Poor	Fair	•	•	•	•	•	•	•
	Dichter <i>et al.</i> (2014)	•	Fair	•	•	•	•	•	•	•
	Ettema <i>et al.</i> (2007a)	•	•	•	•	•	Fair	•	•	•

Table 2. Continued

QOL INSTRUMENT	AUTHOR(S)	INTERNAL CONSISTENCY	RELIABILITY	MEASUREMENT ERROR	CONTENT VALIDITY	STRUCTURAL VALIDITY	HYPOTHESIS TESTING	CROSS-CULTURAL VALIDITY	CRITERION VALIDITY	RESPONSIVENESS
QUALID	Garre-Olmo <i>et al.</i> (2010)	Good	Fair	Fair	•	Fair	Fair	Poor	Poor	•
	Mjorud <i>et al.</i> (2014)	Good	•	•	•	Excellent	•	•	•	•
	Clare <i>et al.</i> (2014a)	Fair	•	•	•	Fair	•	•	•	•
	Falk <i>et al.</i> (2007)	Fair	Poor	•	•	Fair	Poor	Poor	Poor	Poor
	Røen <i>et al.</i> (2015)	Poor	Fair	•	•	•	Poor	Poor	•	•
	Weiner <i>et al.</i> (1999)	Poor	Poor	•	•	Poor	Fair	•	•	•
ADRQL	Edelman <i>et al.</i> (2005)	Poor	•	•	•	Poor	Poor	•	•	•
	Sloane <i>et al.</i> (2005)	Poor	Fair	•	•	•	Poor	•	•	•
	Kasper <i>et al.</i> (2009)	•	•	•	•	•	•	•	•	•
ADRQL revised	Kasper <i>et al.</i> (2009)	Poor	•	•	•	•	•	•	•	•
DCM	Villar <i>et al.</i> (2015)	Good	Poor	•	•	Fair	Fair	•	•	•
	Fossey <i>et al.</i> (2002)	Poor	Fair	•	•	Poor	Poor	•	•	•
	Sloane <i>et al.</i> (2007)	•	Fair	•	•	•	•	•	•	•
	Sloane <i>et al.</i> (2005)	•	Fair	•	•	•	Poor	•	•	•
	Edelman <i>et al.</i> (2005)	•	•	•	•	•	Poor	•	•	•
QOL-AD	León-Salas <i>et al.</i> (2011)	Poor	Poor	Fair	•	•	Fair	•	•	•
QOL-D	Sloane <i>et al.</i> (2005)	Poor	Fair	•	•	•	Poor	•	•	•

reliability scores from a larger number of studies (0.69–0.95).

Measurement error

QUALID and QOL-AD were the only instruments to have measurement error assessed; both were given fair quality ratings.

Content validity

Content validity was assessed in QUALIDEM only, and was rated as fair quality. Structural validity was assessed in six instruments; the instruments most assessed were QUALID, rated in five studies with poor to excellent ratings, and QUALIDEM, rated in four studies with fair to excellent ratings. DQoL and DCM were rated in two studies with fair to poor ratings for both. QOL-AD NH and ADRQL were assessed once with poor ratings.

Hypothesis testing

Hypothesis testing was carried out for eight instruments. The most assessed instruments were QUALID and DCM, which were assessed in four studies with poor to fair ratings. DQoL and QOL-AD NH were assessed in three studies with poor to fair ratings each. ADRQL was assessed in two studies with poor ratings, QUALIDEM and QOL-AD were assessed once with fair ratings, QOL-D had one poor rating.

Cross-cultural validity

Cross-cultural validity was assessed for three instruments (QUALID, QOL-AD NH, and QUALIDEM). QUALID and QOL-AD NH were both assessed twice. QUALID had poor ratings, QOL-AD NH had a poor and a fair rating. QUALIDEM was assessed once and had a poor rating.

Criterion validity

Criterion validity was assessed for two instruments. QUALID was assessed twice, both with poor ratings, and DQoL was assessed once with a poor rating. Only the QUALID was assessed for responsiveness and this had a poor rating.

Feasibility properties

Feasibility properties were extracted for all instruments, except for the case of the QoL-D, in which the original questionnaire and associated materials could not be identified even after contacting the original author. Full details of availability and feasibility properties are presented in Table 3.

Dedicated website

Four of the instruments (QOL-AD, QUALIDEM, ADRQL revised, and DQoL) had dedicated websites where they could be accessed. The ADRQL revised and QOL-AD were both available via the online MAPI trust repository. All other instruments required contacting the original author to access.

User guide

A user guide was accessible for six of the instruments (QOL-AD, QOL-AD NH, QUALIDEM, DCM, ADEQL, ADRQL revised), with many also including additional instructions on the actual questionnaire. We were unable to identify any user instructions for the DQoL and QUALID.

Cost

All instruments were free to access for non-funded research. Additional charges applied for funded research and commercial users in the case of the ADRQL, ADRQL revised, QOL-AD and QOL-AD NH. A single instrument (DCM) was free to use, but required users to attend training, which had associated costs.

Training

Four instruments (DQoL, QOL-AD, QOL-AD NH, and QUALIDEM) did not require any formal training before use of the instrument, whilst the ADRQL and ADRQL revised recommended users to watch a free training video. Only the DCM instrument required users to attend a three-day training course. It was unclear whether the QUALID required training.

Time to complete

Time to complete the instrument ranged from 5 minutes (QUALIDEM, QOL-AD NH) to up to 6 hours (DCM).

Time period to assess

Three instruments captured “present” QoL (QOL-AD, QOL-AD NH, DCM), whilst the remaining instruments assessed QoL over the previous 1 or 2 weeks. The only exception is the DQoL which does not state the time period in which participants should be assessed.

Specialist software

No instrument appeared to require the use of specialist software for the administration, analysis, or interpretation of the instruments.

Table 3. Availability and feasibility properties of identified instruments

SCALE	AVAILABILITY				USABILITY			
	DEDICATED WEBSITE	USER GUIDE	COST	TRAINING REQUIRED	TIME TO COMPLETE	TIME PERIOD TO ASSESS	SPECIALIST SOFTWARE REQUIRED?	SCORING GUIDE
DQoL	No	No	Free	None	10 minutes	Not stated	Not stated	Scoring instructions in original paper (Brod <i>et al.</i> 1999)
QOL-AD	No [*]	Yes, and instructions on instrument	Free to non-funded academic researchers ^a	None	5 mins proxy10–15 mins self-report	Present time (e.g. within the past few weeks)	No	Yes
QOL-AD NH	No	Yes, and instructions on instrument	Free to non-funded academic researchers ^a	None	5 mins proxy10–15 mins self-report	Present time (e.g. within the past few weeks)	No	Yes
QUALIDEM	No	Yes	Free	None	10 minutes	2 weeks Dutch version, 1 week English and German versions	No	Yes, on instrument and in user guide
QUALID	No	No	Free	Not known	5 minutes	1 week	No	Yes, scoring instructions on instrument
DCM	Yes	Yes	Free after training	Required 3 day training (£975)	Up to six hours observation	Present time	Not required but available	Yes
ADRQL	Yes	Yes, and instructions on instrument	Free to non-funded academic researchers ^b	Recommended use of ADRQL training video (free)	10–15 mins	2 weeks	No	Yes
ADRQL Revised	Yes [*]	Yes, and instructions on instrument	Free to non-funded academic researchers ^b	Recommended use of the ADRQL training video (free)	10–15 mins	2 weeks	No	Yes
QoL-D	No	Not known	Not known	Not known	Not known	Not known	Not known	Not known

All information correct as of January 2018.

^{*}Not a dedicated website but available from MAPI trust repository.

^aNon-funded academic researchers - free; Funded academic researchers - €300 per study and €50 per language; Commercial users – Royalty fees €1000 per study and €50 per language. Distribution fees €1000 per study and €50 per language.

^bNon-funded academic researchers — free; Funded academic researchers — €300 per study and €50 per language; Commercial users – Royalty fees 10,000 USD per study and 500 USD per language. Distribution fees €700 per study and €300 per language.

Scoring guide

All instruments had accessible scoring instructions.

Discussion

This study aimed to add usefully to existing reviews of the literature (Bowling *et al.*, 2015; Ready and Ott, 2003). Aspden *et al.* (2014) by assessing the usability properties of instruments in care homes in addition to psychometric properties and study quality. This provides broader information on the pragmatic use of dementia-specific QoL instruments in care homes, which is likely to be of value to researchers, clinicians, and practitioners when selecting instruments for use both in research and in normal practice.

Twenty-five studies were identified that assessed nine dementia specific QoL instruments for use in care homes. This review highlights that even though QoL instruments exist for use in care homes, they may not be accessible or feasible for use. There was limited information about the psychometric properties of most instruments; as a consequence many elements were not assessed. COSMIN scores that were assessed had relatively low ratings with the majority of ratings being poor or fair. The instruments with the most psychometric evidence were QUALID and QUALIDEM. The QUALID instrument had the most extensive evaluation of the two, with seven domains assessed, however, QUALIDEM had better ratings for most of the assessed properties with more excellent and good ratings compared to QUALID. The QUALIDEM has previously been suggested as the best QoL instrument to use for people with dementia in care homes due to the comprehensive assessment of its measurement properties (Aspden *et al.*, 2014).

The QOL-AD instrument is one of the most widely used QoL instruments due to its good psychometric properties and apparent ease of use (Bowling *et al.*, 2015). However, findings from the review show limited use of the QOL-AD in care settings, with the original QOL-AD only identified in one study, and the nursing home version identified in six studies. The QOL-AD NH instrument, which is adapted from QOL-AD was assessed in six studies; however, the ratings were mainly poor or fair. These findings are in line with a previous review which found that the QOL-AD NH properties were poor because of small sample sizes and methodological quality of the studies (Aspden *et al.*, 2014).

It is widely accepted that measuring QoL in care settings is important. This is usually carried out as a part of research with the focus being to understand changes to QoL over time or to assess outcomes of

interventions (Clare *et al.*, 2014b; Hoe *et al.*, 2009). There is, however, an additional need and potential benefit in understanding the QoL of care home residents outside of research, in routine care practice. The lack of routine measurement of QoL in care homes will be in part due to differences in the current instruments and a lack of consensus about what a QoL instrument should contain. The frameworks of QoL in dementia underpinning each instrument influence and shape the instrument and its content (Missotten *et al.*, 2016). The instruments discussed here are based on a small number of models. Most are based solely, or in part, on the work of Lawton (Missotten *et al.*, 2016), which states that QoL in dementia is multidimensional and consists of both objective and subjective components. A health-related QoL definition, which only includes aspects of QoL affected by a health condition, and the adaptation-coping model, which is concerned with adaptation to the consequences of dementia (Dröes *et al.*, 2010) have also been used. However, it is often not clear what conceptual framework instruments are based on, and what assumptions are made (Bowling *et al.*, 2015; Missotten *et al.*, 2016). This may dissuade some from using QoL instruments, and the differences in the content of instruments may influence whether or not specific instruments are used. This, coupled with what it is that practitioners, researchers, or clinicians want to gain from measuring QoL will influence the decision about what instrument to use. However a fundamental problem is likely to be limited access to and experience of QoL instruments as well as a lack of requirement by regulators to use such approaches. Most are difficult to find and access online; often requiring contacting the original author or registering with online instrument repositories to request access to specific instruments. This will be difficult for care staff who may not be aware of the different instruments available and may not have the skills and resources needed to identify them.

Routine use of QoL instruments might be of value at an individual resident level and also at an aggregate level providing insights into the QoL of residents as a whole and indirectly the home as a whole. QoL measurement has the potential to improve the provision of person-centered or holistic care by encouraging care staff to focus more on the individual and less on impairments of function (Edvardsson *et al.*, 2014). The inclusion of an instrument to specifically measure QoL in normal care practice, and used in conjunction with existing care practices, procedures, and documentation, could enhance the opportunity to improve care quality by providing more person-centered information on residents and providing feedback on the effect of changes over time. In the US, a comprehensive assessment of a variety of care

features is completed for residents in long-term care facilities. The Resident Assessment Instrument (RAI-NH) is completed periodically to collect and record relevant information about resident care. The aim of its use is to inform holistic care and ensure good QoL and quality of care. Implementation and use of the RAI-NH has been shown to be associated with improvements in resident outcomes (Fries *et al.*, 1997; Phillips *et al.*, 1997). The integration of a QoL instrument into such a system and used regularly could enhance the assessment of resident outcomes by providing complementary information and therefore a broader understanding of wellbeing of the resident and when aggregated further insights into the quality of the care home. In the UK and many other countries there is no such mandated combined assessment schedule in the care sector and no culture of routine measurement; resident and care factors are documented by staff, in a less systematic manner than in the RAI-NH. Therefore, integrating QoL measurement into routine care practice would need a different approach. Regardless of the systems that currently exist in homes, such action would require appropriate instruments that are accessible for and usable by care home staff. Of all the instruments that were available, the QOL-AD, QUALIDEM, and ADLQOL revised were accessible via simple internet searches and had a dedicated web page. Importantly, many of the measures were difficult to access, and required contacting the original author (e.g., DQoL). Overall, the QUALIDEM was assessed as the most accessible; being free to use, with all information regarding administration and scoring available in an extensive user guide. However, although an English language version of QUALIDEM is available, it has not been validated in an English sample as far as could be determined from this review. Therefore, despite QUALIDEM being the most psychometrically robust and accessible instrument there is still some concern regarding whether it is suitable for use in an English-speaking sample and setting.

In this review we focus on dementia-specific QoL instruments. A previous review by Aspden and colleagues suggested that one instrument should be used for people with dementia and another for people without dementia (Aspden *et al.*, 2014). We understand the clarity of purpose behind this, but believe that in routine care practice, when those with dementia are in the majority, it may be preferable to use a single dementia-specific instrument for three main reasons. First, using two different instruments makes it difficult to aggregate scores for homes overall, and changes to resident status (from no diagnosis to diagnosis) would require a change in instrument, thus meaning a lack of consistent measurement for that individual resident over time. Second, dual measurement is dependent on an

accurate diagnosis of a dementia having been made for each resident and this is unlikely to be the case. Diagnosis rates in the UK are amongst the highest in the world but are still only around 67% (Department of Health, 2016). Many without a formal diagnosis of dementia in care homes will in fact have dementia, and evidence of high prevalence of dementia in care homes mean that most residents would be appropriate for the use of a dementia-specific QoL instrument (Jagger and Lindesay, 1997; Livingston *et al.*, 2017). Finally, the questions contained in QoL instruments need to be appropriate for people with dementia and for use in care settings. Few generic measures of QoL have been developed specifically for use in care home settings, and very few have been validated in this setting. They can, therefore, contain inappropriate questions, often reflecting the opportunity to perform a function rather than the ability to perform it (Hall *et al.*, 2011). We believe that the content of dementia-specific QoL instruments may be a better fit for care home residents, even those without dementia, than generic instruments. This can be posed as an empirical question and requires research to be done to test this hypothesis. On balance we believe a strategy of dual measurement may introduce more measurement error than it prevents with the addition of unnecessary complexity.

Despite differences in the content of instruments and the findings presented in this review, the use of existing QoL instruments is preferable to using no QoL instruments at all. They provide potentially valuable information compared with not using them. There is, however, a need for the development of instruments that are developed and/or validated to measure QoL in care home settings and that are accessible, easy to use, and contain appropriate questions. The future development and adaptation of instruments needs to consider these care-home specific points to create instruments that are of demonstrable benefit and lead to the wider use of QoL instruments in care homes.

In this review we do not recommend the use of one instrument as being the best or most appropriate to use. Instead, because there is no gold standard instrument, we argue that there are factors other than psychometric properties that should be considered when deciding on an appropriate instrument to use in care settings, particularly if the care staff will be involved in collecting data. We conclude that new instruments need to be developed either from existing measures or de novo that are specific to the care home setting, that have acceptable psychometric properties and that are readily accessible and usable in these settings. Deciding on what instrument is the most useful in care settings is difficult given the differences in care homes themselves.

Issues of time and resources and the benefit that accrues from their use will always be to the fore, so data on these parameters are needed. The development or adaptation of instruments that can be used in care homes by both researchers and by care staff routinely in normal care practice would be useful. If adapted, the content of such instruments would need to remain largely the same to maintain reliability and validity of the instrument. However, the format or layout of the instrument questions, and user guides or instructions would need to be adapted for use in care homes and by care staff. Researchers will be specifically trained and experienced in using standardised instruments and are likely to have more time to complete instruments. Care staff are likely to need more specific guidelines to follow and a layout that reduces the risk of recording error and missing data.

It is important to highlight several limitations of this review. First, this review did not include all measures of dementia-specific QoL, because they did not meet inclusion criteria of this review. For example, the Philadelphia Geriatric Centre Affect Rating Scale (PGC-ARS) (Lawton *et al.*, 1996) and the Resident and Staff Observation Checklist Quality of Life measure (RSOC-QOL) (Sloane *et al.*, 1991) were not included in this review despite being included in a review by Aspden and colleagues as the instruments were not assessed to be measures of QoL (Aspden *et al.*, 2014). Second, there is no established means of assessing the usability of instruments in care homes, and therefore interpretation of what is considered “accessible” or “practical” may differ depending on the user. Third, this review focused on disease-specific instruments of QoL and makes the assumption that the majority of residents in a care home would have dementia; this may not be the case and could affect the validity of instruments. Finally, this review only included articles published in English, meaning that non-English instruments may not have been captured.

Conclusion

The number of high quality easily accessible instruments available for use in care homes is low. In general, the psychometric analysis of instruments revealed them to function poorly in care home settings with a limited number of psychometric elements assessed for each instrument in any single study. Furthermore, the quality ratings of each instrument were weak with very few receiving excellent or good ratings. Instruments with the best psychometric assessments were not necessarily easily accessible, and those that were readily available had some of the poorest quality and most limited

assessment. The findings of this review indicate that there is a need for further large and well-designed evaluations of the properties of these instruments when used in care homes, including head-to-head comparisons. Further work is also needed in order to ensure that any instruments are made easily available for use by care staff if they are to be used in a non-research or clinical capacity. Due to poor psychometric and feasibility properties, none of the instruments currently available seem suited to routine use by care staff in care homes. Further development of existing instruments or the generation of new methods for the measurement of QoL in care homes would be of potential value given the need to assure and improve the quality of care of residents in care homes with dementia.

Conflict of interest

None.

Description of authors' roles

LH completed the search, assessed the articles, and wrote the first draft of the review. NF and TEP assisted with article assessment and screening, and provided inputs and amendments to the review. NT and SB provided input and amendments throughout the development of the review.

Supplementary material

To view supplementary material for this article, please visit <https://doi.org/10.1017/S1041610218002259>.

References

- Adler, E. and Resnick, B. (2010). Reliability and validity of the dementia quality of life measure in nursing home residents. *Western Journal of Nursing Research*, 32, 686–704.
- Albert, S. M. *et al.* (1996). Quality of life in patients with Alzheimer's disease as reported by patient proxies. *Journal of the American Geriatrics Society*, 44, 1342–1347.
- Aspden, T., Bradshaw, S. A., Playford, E. D. and Riazi, A. (2014). Quality-of-life measures for use within care homes: a systematic review of their measurement properties. *Age and Ageing*, 43, 596–603.
- Beer, C., Bosboom, P., Almeida, O. P. and Flicker, L. (2009). Rating the quality of life of people with dementia living in residential care facilities in routine research practice. *Age and Ageing*, 38, 343–346.
- Black, N. (2013). Patient reported outcome measures could help transform healthcare. *BMJ*, 346, 1–5.

- Bouman, A. I. E., Ettema, T. P., Wetzels, R. B., van Beek, A. P. A., de Lange, J. and Dröes, R. M.** (2011). Evaluation of Qualidem: a dementia-specific quality of life instrument for persons with dementia in residential settings; scalability and reliability of subscales in four Dutch field surveys. *International Journal of Geriatric Psychiatry*, 26, 711–722.
- Bowling, A. *et al.*** (2015). Quality of life in dementia: a systematically conducted narrative review of dementia-specific measurement scales. *Aging & Mental Health*, 19, 13–31.
- Bradford Dementia Group.** (2005). *DCM 8 User's Manual*. Bradford: University of Bradford.
- Brod, M., Stewart, A. L., Sands, L. and Walton, P.** (1999). Conceptualization and measurement of quality of life in dementia: the dementia quality of life instrument (DQoL). *The Gerontologist*, 39, 25–36.
- Care Quality Commission.** (2017a). About us. Available at: <http://www.cqc.org.uk/what-we-do/services-we-regulate/care-homes>.
- Care Quality Commission.** (2017b). The state of adult social care services 2014 to 2017. Available at: <http://www.cqc.org.uk/publications/major-report/state-adult-social-care-services-2014-2017>.
- Centre for Disease Control and Prevention.** (2017). FastStats. Available at: <https://www.cdc.gov/nchs/fastats/nursing-home-care.htm>; last accessed 25 April 2018.
- Clare, L., Quinn, C., Hoare, Z., Whitaker, R. and Woods, R. T.** (2014a). Care staff and family member perspectives on quality of life in people with very severe dementia in long-term care: a cross-sectional study. *Health and Quality of Life Outcomes*, 12, 175.
- Clare, L. *et al.*** (2014b). Trajectories of quality of life in early-stage dementia: individual variations and predictors of change. *International Journal of Geriatric Psychiatry*, 29, 616–623.
- Crespo, M., de Quirós, M. B., Gómez, M. M. and Hornillos, C.** (2011). Quality of life of nursing home residents with dementia: a comparison of perspectives of residents, family, and staff. *The Gerontologist*, 52, gnr080.
- Department of Health.** (2016). Prime Minister's challenge on dementia 2020. Available at: <https://www.gov.uk/government/publications/challenge-on-dementia-2020-implementation-plan>.
- Department of Health, Prime Minister's Office.** (2015). Prime Minister's challenge on dementia. Available at: <https://www.gov.uk/government/publications/prime-ministers-challenge-on-dementia-2020>.
- Dichter, M., Bartholomeyczik, S., Nordheim, J., Achterberg, W. and Halek, M.** (2011). Validity, reliability, and feasibility of a quality of life questionnaire for people with dementia. *Zeitschrift für Gerontologie und Geriatrie*, 44, 405–410.
- Dichter, M., Halek, M., Dortmann, O., Meyer, G. and Bartholomeyczik, S.** (2013). Measuring the quality of life of people with dementia in nursing homes in Germany - the study protocol for the QoL-Dem Project. *Psycho-Social Medicine*, 10, Doc06.
- Dichter, M., Schwab, C. G. G., Meyer, G., Bartholomeyczik, S., Dortmann, O. and Halek, M.** (2014). Measuring the quality of life in mild to very severe dementia: testing the inter-rater and intra-rater reliability of the German version of the QUALIDEM. *International Psychogeriatrics*, 26, 825–836.
- Dichter, M., Schwab, C. G. G., Meyer, G., Bartholomeyczik, S. and Halek, M.** (2016a). Item distribution, internal consistency and inter-rater reliability of the German version of the QUALIDEM for people with mild to severe and very severe dementia. *BMC Geriatrics*, 16, 126.
- Dichter, M., Wolschon, E.-M., Meyer, G. and Köpke, S.** (2016b). Cross-cultural adaptation of the German version of the Quality of Life in Alzheimer's Disease scale - Nursing Home version (QoL-AD NH). *International Psychogeriatrics*, 28, 1–2.
- Dröes, R. M., van Mierlo, L. D., van der Roest, H. G. and Meiland, F. J. M.** (2010). Focus and effectiveness of psychosocial interventions for people with dementia in institutional care settings from the perspective of coping with the disease. *Non-Pharmacological Therapies in Dementia*, 1, 23.
- Edelman, P., Fulton, B. R., Kuhn, D. and Chang, C.-H.** (2005). A comparison of three methods of measuring dementia-specific quality of life: perspectives of residents, staff, and observers. *The Gerontologist*, 45, 27–36.
- Edvardsson, D., Sandman, P. O. and Borell, L.** (2014). Implementing national guidelines for person-centered care of people with dementia in residential aged care: effects on perceived person-centeredness, staff strain, and stress of conscience. *International Psychogeriatrics*, 26, 1171–1179.
- Ettema, T. P., Dröes, R.-M., de Lange, J., Mellenbergh, G. J. and Ribbe, M. W.** (2007a). QUALIDEM: development and evaluation of a dementia specific quality of life instrument. Scalability, reliability and internal structure. *International Journal of Geriatric Psychiatry*, 22, 549–556.
- Ettema, T. P., Dröes, R.-M., de Lange, J., Mellenbergh, G. J. and Ribbe, M. W.** (2007b). QUALIDEM: development and evaluation of a dementia specific quality of life instrument—validation. *International Journal of Geriatric Psychiatry*, 22, 424–430.
- Falk, H., Persson, L.-O. and Wijk, H.** (2007). A psychometric evaluation of a Swedish version of the Quality of Life in Late-Stage Dementia (QUALID) scale. *International Psychogeriatrics*, 19, 1040–1050.
- Folstein, M. F., Folstein, S. E. and McHugh, P. R.** (1975). "Mini-mental state": a practical method for grading the cognitive state of patients for the clinician. *Journal of Psychiatric Research*, 12, 189–198.
- Fossey, J., Lee, L. and Ballard, C.** (2002). Dementia care mapping as a research tool for measuring quality of life in care settings: psychometric properties. *International Journal of Geriatric Psychiatry*, 17, 1064–1070.
- Fries, B. E., Hawes, C., Morris, J. N., Phillips, C. D., Mor, V. and Park, P. S.** (1997). Effect of the national resident assessment instrument on selected health conditions and problems. *Journal of the American Geriatrics Society*, 45, 994–1001.
- Garre-Olmo, J. *et al.*** (2010). Cross-cultural adaptation and psychometric validation of a Spanish version of the Quality of Life in Late-Stage Dementia Scale. *Quality of Life Research*, 19, 445–453.
- Hall, S., Opio, D., Dodd, R. H. and Higginson, I. J.** (2011). Assessing quality-of-life in older people in care homes. *Age and Ageing*, 40, 507–512.

- Hoe, J., Hancock, G., Livingston, G., Woods, B., Challis, D. and Orrell, M.** (2009). Changes in the quality of life of people with dementia living in care homes. *Alzheimer Disease and Associated Disorders*, 23, 285–290.
- Jagger, C. and Lindesay, J.** (1997). Residential care for elderly people: the prevalence of cognitive impairment and behavioural problems. *Age and Ageing*, 26, 475–480.
- Kasper, J. D., Black, B. S., Shore, A. D. and Rabins, P. V.** (2009). Evaluation of the validity and reliability of the Alzheimer disease-related quality of life assessment instrument. *Alzheimer Disease and Associated Disorders*, 23, 275–284.
- Lawton, M. P., Haitsma, K. V. and Klapper, J.** (1996). Observed affect in nursing home residents with Alzheimer's disease. *The Journals of Gerontology Series B: Psychological Sciences and Social Sciences*, 51B, P3–P14.
- León-Salas, B., Logsdon, R. G., Olazarán, J. and Martínez-Martín, P.** (2011). Psychometric properties of the Spanish QoL-AD with institutionalized dementia patients and their family caregivers in Spain. *Aging & Mental Health*, 15, 775–783.
- Livingston, G. et al.** (2017). Prevalence of and associations with agitation in residents with dementia living in care homes: MARQUE cross-sectional study. *British Journal of Psychiatry Open*, 3, 171–178.
- Logsdon, R. G., Gibbons, L. E., McCurry, S. M. and Teri, L.** (1999). Quality of life in Alzheimer's disease. *Journal of Mental Health and Aging*, 5, 21–32.
- Missotten, P., Dupuis, G. and Adam, S.** (2016). Dementia-specific quality of life instruments: a conceptual analysis. *International Psychogeriatrics*, 28, 1245–1262.
- Mjorud, M. et al.** (2014). Principal component analysis of the Norwegian version of the Quality of Life in Late-Stage Dementia Scale. *Dementia and Geriatric Cognitive Disorders*, 37, 265–275.
- Mokkink, L. B. et al.** (2010). The COSMIN study reached international consensus on taxonomy, terminology, and definitions of measurement properties for health-related patient-reported outcomes. *Journal of Clinical Epidemiology*, 63, 737–745.
- Morris, J. C.** (1997). Clinical dementia rating: a reliable and valid diagnostic and staging measure for dementia of the Alzheimer type. *International Psychogeriatrics*, 9, 173–176.
- Moyle, W., Gracia, N., Murfield, J. E., Griffiths, S. G. and Venturato, L.** (2012). Assessing quality of life of older people with dementia in long-term care: a comparison of two self-report measures. *Journal of Clinical Nursing*, 21, 1632–1640.
- NIHR.** (2016). Understanding care homes. Available at: <http://enrich.nihr.ac.uk/page/understanding-care-homes>; last accessed 17 September 2017.
- Phillips, C. D. et al.** (1997). Association of the resident assessment instrument (RAI) with changes in function, cognition, and psychosocial status. *Journal of the American Geriatrics Society*, 45, 986–993.
- Prince, M. et al.** (2014). Dementia UK: update. Available at: https://www.alzheimers.org.uk/sites/default/files/migrate/downloads/dementia_uk_update.pdf; last accessed 11 July 2017.
- Rabins, P., Kasper, J. D., Kleinman, L., Black, B. S. and Patrick, D. L.** (1999). Concepts and methods in the development of the ADRQL: an instrument for assessing health-related quality of life in persons with Alzheimer's disease. *Journal of Mental Health and Aging*, 5, 33–48.
- Ready, R. E. and Ott, B. R.** (2003). Quality of life measures for dementia. *Health and Quality of Life Outcomes*, 1, 11.
- Reisberg, B.** (1988). Functional assessment staging (FAST). *Psychopharmacology Bulletin*, 24, 653–659.
- Reisberg, B., Ferris, S. H., de Leon, M. J. and Crook, T.** (1982). The Global Deterioration Scale for assessment of primary degenerative dementia. *The American Journal of Psychiatry*, 139, 1136–1139.
- Røen, I., Selbaek, G., Kirkevold, Ø., Engedal, K., Lerdal, A. and Bergh, S.** (2015). The reliability and validity of the Norwegian version of the Quality of Life in Late-Stage Dementia Scale. *Dementia and Geriatric Cognitive Disorders*, 40, 233–242.
- Sloane, P. D., Mathew, L. J., Scarborough, M., Desai, J. R., Koch, G. G. and Tangen, C.** (1991). Physical and pharmacologic restraint of nursing home patients with dementia: impact of specialized units. *JAMA*, 265, 1278–1282.
- Sloane, P. D., Zimmerman, S., Williams, C. S., Reed, P. S., Gill, K. S. and Preisser, J. S.** (2005). Evaluating the quality of life of long-term care residents with dementia. *The Gerontologist*, 45, 37–49.
- Sloane, P. D. et al.** (2007). Dementia care mapping as a research tool. *International Journal of Geriatric Psychiatry*, 22, 580–589.
- Smith, S. et al.** (2005). Measurement of health-related quality of life for people with dementia: development of a new instrument (DEMQOL) and an evaluation of current methodology. *Health Technology Assessment*, 9, 1–93.
- Villar, F., Vila-Miravent, J., Celdrán, M. and Fernández, E.** (2015). Assessing the well-being of patients with dementia: validity and reliability of the Spanish version of the Dementia Care Mapping (DCM) tool / La evaluación del bienestar en pacientes con demencia: validez y fiabilidad de la versión española del Dementia Care Mapping (DCM). *Estudios de Psicología*, 36, 389–412.
- Weiner, M., Martin-Cook, K., Svetlik, D. A., Saine, K., Foster, B. and Fontaine, C. S.** (1999). The quality of life in late-stage dementia (QUALID) scale. *Journal of the American Medical Disorders Association*, 1, 114–116.